

monitored for scrapie disease until the flock, and all first generation progeny resulting from the semen imported in accordance with this section, qualifies as a “Certified” flock.

(f) Except for sheep semen being placed in Certifiable Class C flocks, the certificate accompanying the sheep semen imported under paragraph (a) of this section must contain the following statement: “The semen identified on this certificate has been collected from a sire that has been monitored by a salaried veterinary officer of [*name of region of origin*], for [*number of months*], in the same source flock which had been determined by the Administrator, APHIS, prior to the exportation of the semen to the United States, to be equivalent to [*certification level*] of the Voluntary Scrapie Flock Certification Program authorized under 9 CFR part 54, subpart B.”

(1) The Administrator will determine, based upon information supplied by the importer, whether the donor animal’s flock participates in a program in the region of origin that is equivalent to the Voluntary Scrapie Flock Certification Program, and if so, at what level the source flock would be classified.

(2) In order for the Administrator to make a determination, the importer must supply the following information with the application for an import permit, no less than 1 month prior to the anticipated date of importation:

(i) The name, title, and address of a knowledgeable official in the veterinary services of the region of origin;

(ii) The details of scrapie control programs in the region of origin, including information on disease surveillance and border control activities and the length of time these activities have been in effect;

(iii) Any available information concerning additions, within the 5 years immediately preceding collection of the semen, to the flock of the semen donor;

(iv) Any available data concerning disease incidence, within the 5 years immediately preceding collection of the semen in the donor animal’s flock, including, but not limited to, the results of diagnostic tests, especially

histopathology tests, conducted on any animals in the flock;

(v) Information concerning the health, within the 5 years immediately preceding collection of the semen, of other ruminants, flocks, and herds with which the donor animal and the donor animal’s flock might have had physical contact, and a description of the type and frequency of the physical contact; and

(vi) Any other information requested by the Administrator in specific cases as needed to make a determination.

(g) All first generation progeny resulting from semen imported under this section are subject to the requirements of 9 CFR part 54 and all other applicable regulations.

(Approved by the Office of Management and Budget under control numbers 0579–0040 and 0579–0101)

[61 FR 17242, Apr. 19, 1996, as amended at 62 FR 56026, Oct. 28, 1997; 64 FR 23179, April 30, 1999]

§ 98.38 Restrictions on the importation of swine semen from parts of the European Union.

In addition to meeting all other applicable provisions of this part, swine semen imported from the region of the European Union consisting of Austria, Belgium, Germany (except for the Kreis Uckermark in the Land of Brandenburg; the Kreis Oldenburg, the Kreis Soltau-Fallingb., and the Kreis Vechta in the Land of Lower Saxony; the Kreis Heinsberg and the Kreis Warendorf in the Land of Northrhine-Westphalia; the Kreis Bernkastel-Wittlich, the Kreis Bitburg-Prüm, the Kreis Donnersbergkreis, the Kreis Rhein-Hunsrück, the Kreis Südliche Weinstraße, and the Kreis Trier-Saarburg in the Land of Rhineland Palatinate; and the Kreis Altmarkkreis in the Land of Saxony-Anhalt); Greece, Italy (except for the Regions of Emilia-Romagna, Piemonte, and Sardegna), the Netherlands, and Portugal must meet the following conditions:

(a) The semen must come only from a semen collection center approved for export by the veterinary services of the national government of the country of origin;

(b) The donor boar must not have lived in a region when the region was

classified in § 94.10(a) as one in which classical swine fever is known to exist, and must not have transited such a region unless moved directly through the region in a sealed means of conveyance with the seal determined to be intact upon arrival at the point of destination;

(c) The donor boar must never have been commingled with swine that have been in a region when the region was classified in § 94.10(a) as one in which classical swine fever is known to exist;

(d) The donor boar must be held in isolation for at least 30 days prior to entering the semen collection center;

(e) No more than 30 days prior to being held in isolation as required by paragraph (d) of this section, the donor boar must be tested with negative results with a classical swine fever test approved by the Office International des Epizooties;

(f) No equipment or materials used in transporting the donor boar from the farm of origin to the semen collection center may have been used previously for transporting swine that do not meet the requirements of this section, unless such equipment or materials has first been cleaned and disinfected;

(g) The donor boar must be observed at the semen collection center by the center veterinarian, and exhibit no clinical signs of classical swine fever;

(h) Before the semen is exported to the United States, the donor boar must be held at the semen collection center for at least 40 days following collection of the semen, and, along with all other swine at the semen collection center, exhibit no clinical signs of classical swine fever; and

(i) The semen must be accompanied to the United States by a certificate issued by a salaried veterinary officer of the national government of the country of origin, stating that the provisions of paragraphs (a) through (h) of this section have been met.³

(Approved by the Office of Management and Budget under control number 0579-0218)

[68 FR 16940, Apr. 7, 2003]

³The certification required may be placed on the certificate required under § 98.35(c) or may be contained in a separate document.

PART 99—RULES OF PRACTICE GOVERNING PROCEEDINGS UNDER CERTAIN ACTS

Subpart A—General

Sec.

99.1 Scope and applicability of rules of practice.

Subpart B—Supplemental Rules of Practice

99.10 Stipulations.

AUTHORITY: 7 U.S.C. 8301-8317; 7 CFR 2.22, 2.80, and 371.4.

SOURCE: 48 FR 30095, June 30, 1983, unless otherwise noted. Redesignated at 52 FR 29502, Aug. 10, 1987.

Subpart A—General

§ 99.1 Scope and applicability of rules of practice.

The Uniform Rules of Practice for the Department of Agriculture promulgated in subpart H of part 1, subtitle A, title 7, Code of Federal Regulations, are the Rules of Practice applicable to adjudicatory, administrative proceedings under the following statutory provisions:

Act of May 29, 1884, commonly known as the Animal Industry Act, section 7, as amended (21 U.S.C. 117),

Act of August 30, 1890, section 6, as amended (21 U.S.C. 104),

Act of February 2, 1903, commonly known as the Cattle Contagious Diseases Act of 1903, section 3, as amended (21 U.S.C. 122),

Act of July 2, 1962, section 6(a), as amended (21 U.S.C. 134e),

Act of May 6, 1970, section 2, as amended (21 U.S.C. 135a).

The Animal Health Protection Act, section 10414 (7 U.S.C. 8313)

In addition, the Supplemental Rules of Practice set forth in subpart B of this part shall be applicable to such proceedings.

[48 FR 30095, June 30, 1983. Redesignated at 52 FR 29502, Aug. 10, 1987, as amended at 68 FR 6345, Feb. 7, 2003]

Subpart B—Supplemental Rules of Practice

§ 99.10 Stipulations.

(a) At any time prior to the issuance of a complaint seeking a civil penalty